# Food and Drug Administration, HHS

- (iv) The labeling for feed or feed ingredients to which formaldehyde has been added under the provisions of paragraph (b)(1) of this section is required to carry the following statement: "Treated with formaldehyde to maintain feed *Salmonella* negative. Use within 21 days."
- (3) To assure safe use of the additive, in addition to the other information required by the Act, the label and labeling shall contain:
- (i) Appropriate warnings and safety precautions concerning formaldehyde.
- (ii) Statements identifying formaldehyde as a poison with potentials for adverse respiratory effects.
- (iii) Information about emergency aid in case of accidental inhalation.
- (iv) Statements reflecting requirements of applicable sections of the Superfund Amendments and Reauthorization Act (SARA), and the Occupational Safety and Health Administration's (OSHA) human safety guidance regulations.
- (v) Contact address and phone number for reporting adverse reactions or to request a copy of the Materials Safety Data Sheet (MSDS).
- [41 FR 38652, Sept. 10, 1976, as amended at 54 FR 18281, Apr. 28, 1989; 61 FR 15704, Apr. 9, 1996; 63 FR 53580, Oct. 6, 1998; 68 FR 65633, Nov. 21, 2003]

# § 573.480 Formic acid.

The food additive, formic acid, may be safely used in accordance with the following conditions:

- (a) The additive is used as a preservative in hay crop silage in an amount not to exceed 2.25 percent of the silage on a dry weight basis or 0.45 percent when direct cut, as follows:
- (1) The top foot of silage stored should not contain formic acid and
- (2) Silage should not be fed to livestock within 4 weeks of treatment.
- (b) The additive is used or intended for use as a feed acidifying agent, to lower the pH, in complete swine feeds at levels not to exceed 1.2 percent of the complete feed.
- (1) The additive consists of not less than 85 percent formic acid (CAS 64-18-6).
- (2) The additive meets the following specifications:

- (i) Free methyl alcohol not to exceed 1,000 parts per million (ppm);
- (ii) Methyl formate not to exceed 1,000 ppm; and
- (iii) Moisture not to exceed 15 percent.
- (3) To assure safe use of the additive, in addition to the other information required by the Federal Food, Drug and Cosmetic Act, the label and labeling shall contain:
  - (i) The name of the additive.
- (ii) Adequate directions for use including a statement that formic acid must be uniformly applied and thoroughly mixed into complete swine feeds and that the complete swine feeds treated shall be labeled as containing formic acid.
- (4) To assure safe use of the additive, in addition to the other information required by the Federal Food, Drug, and Cosmetic Act and paragraph (b)(3) of this section, the label and labeling shall contain:
- (i) Appropriate warnings and safety precautions concerning formic acid (85 percent formic acid).
- (ii) Statements identifying formic acid (85 percent formic acid) as a corrosive and possible severe irritant.
- (iii) Information about emergency aid in case of accidental exposure.
- (A) Statements reflecting requirements of applicable sections of the Superfund Amendments and Reauthorization Act (SARA), and the Occupational Safety and Health Administration's (OSHA) human safety guidance regulations.
- (B) Contact address and telephone number for reporting adverse reactions or to request a copy of the Material Safety Data Sheet (MSDS).

[76 FR 7106, Feb. 9, 2011]

# § 573.500 Condensed, extracted glutamic acid fermentation product.

Condensed, extracted glutamic acid fermentation product may be safely used in animal feed under the following conditions:

(a) The additive is a concentrated mixture of the liquor remaining from the extraction of glutamic acid, combined with the cells of *Corynebacterium lilium* used to produce the glutamic acid.

# §573.520

- (b) It is used or intended for use as follows:
- (1) In poultry feed as a source of protein in an amount not to exceed 5 percent of the total ration.
- (2) In cattle feed as a source of protein in an amount not to exceed 10 percent of the feed.
- (c) In order to assure safe use, the label and labeling of the additive shall bear, in addition to the other information required by the Act, the following:
  - (1) The name of the additive.
- (2) A statement of the concentration of the additive contained in any mixture.
  - (3) Adequate directions for use.

#### § 573.520 Hemicellulose extract.

Hemicellulose extract may be safely used in animal feed when incorporated therein in accordance with the following conditions:

- (a) The additive is produced from the aqueous extract obtained by the treatment of wood with water at elevated temperatures (325 degrees–535 degrees F) and pressure (80 to 900 pounds per square inch) and contains primarily pentose and hexose sugars.
- (b) The additive may be used in a liquid or dry state with the liquid product containing not less than 55 percent carbohydrate and the dry product containing not less than 84 percent carbohydrate.
- (c) The additive is used as a source of metabolizable energy in animal feed in accordance with good manufacturing and feeding practices.
- [41 FR 38652, Sept. 10, 1976, as amended at 43 FR 11181, Mar. 17, 1978]

# $\S 573.530$ Hydrogenated corn syrup.

- (a) *Identity*. The product is produced by hydrogenation of corn syrup over a nickel catalyst.
- (b) Specifications. The product contains 70 percent hydrogenated corn syrup and a maximum of 0.5 percent reducing sugars.
- (c) *Uses*. The product is used as a humectant and plasticizer in preparation of soft-moist dog and cat foods.
- (d) *Limitations*. The product is preferably stored in a closed, stainless steel or aluminum container. The level of use of the product shall not exceed

15 percent of the total weight of the pet food formulation.

- (e) *Labeling*. The labeling shall bear, in addition to other information required by the Act:
  - (1) The name of the additive.
- (2) Adequate directions for use in accordance with the provisions in paragraph (d) of this section.

[45 FR 22920, Apr. 4, 1980]

# § 573.540 Hydrolyzed leather meal.

- (a) *Identity*. Hydrolyzed leather meal is produced from leather scraps that are treated with steam for not less than 33 minutes at a pressure of not less than 125 pounds per square inch.
- (b) *Specifications*. The additive shall conform to the following percent-by-weight specifications:

Moisture, not less than 5 percent nor more than 10 percent.

Crude protein, not less than 60 percent. Crude fat, not less than 5 percent. Crude fiber, not more than 6 percent. Chromium, not more than 2.75 percent.

- (c) *Use.* It is used or intended for use as a source of protein in swine feeds in an amount not to exceed 1.0 percent by weight of the finished feed.
- (d) *Labeling*. The labels and labeling shall bear, in addition to the other information required by the Act:
- (1) The name of the additive, hydrolyzed leather meal.
- (2) Adequate directions to provide finished feeds complying with paragraph (c) of this section.

### § 573.560 Iron ammonium citrate.

Iron ammonium citrate may be safely used in animal feed in accordance with the following prescribed conditions:

- (a) The additive is the chemical green ferric ammonium citrate.
- (b) The additive is used or intended for use as an anticaking agent in salt for animal consumption so that the level of iron ammonium citrate does not exceed 25 parts per million (0.0025 percent) in the finished salt.
- (c) To assure safe use of the additive the label or labeling of the additive shall bear, in addition to the other information required by the Act:
- (1) The name of the additive.